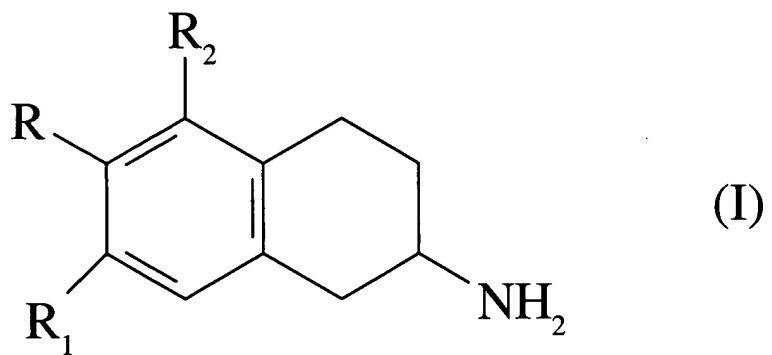


AMENDMENTS TO THE CLAIMS:

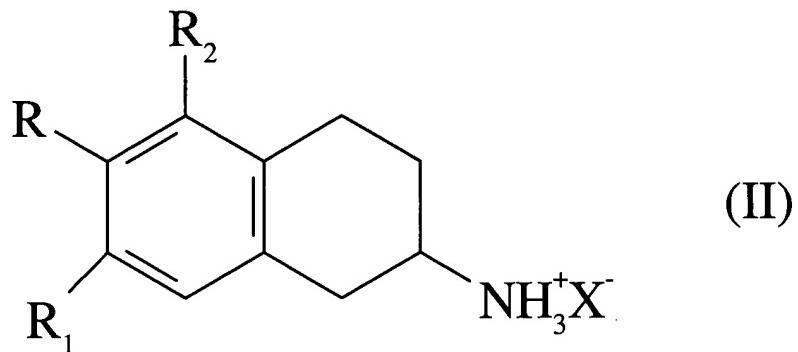
This listing of claims will replace all prior versions, and listings, of claims in the application:

1.-7. Canceled.

8. (Previously Presented) A method of treating an inflammatory and/or autoimmune pathology induced by inflammatory cytokines, which method comprises administering to a patient in need of same an effective amount of a 2-aminotetraline of the formula (I)



or a pharmacologically acceptable salt of the formula (II)



wherein:

R and R₁ are independently halogen, hydroxy, or C₁-C₄ alkoxy optionally substituted in position ω with a group selected from OH, NH₂ or NR₃R₄, wherein R₃ and R₄ are independently H, C₁-C₄ alkyl, unsubstituted or substituted in position ω with groups OH, NH₂, C₁-C₄ alkanoyl, C₁-C₄ alkyl, carbamoyl, carbamoyloxy, amino, or amino-substituted NR₃R₄, where R₃ and R₄ have the above meanings,

R₂ is hydrogen, halogen, hydroxy or methoxy,

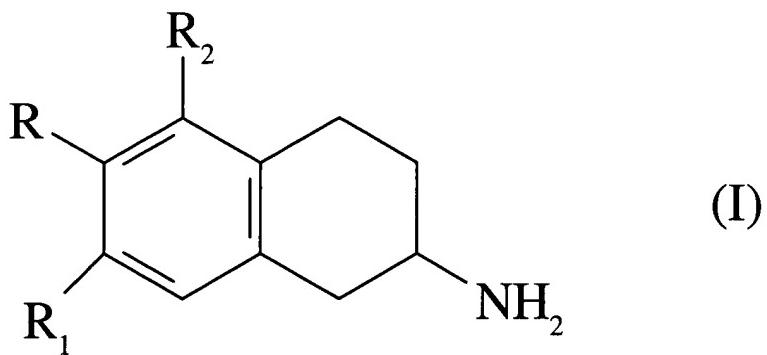
with the proviso that the 2-aminotetraline excludes (a) R=R₁=CH₃O or OH, R₂=H,

(b) R=F, R₁=CH₃O or OH, R₂=H, (c) R₁=-OCH₃, R=CH₃ and R₂=H, or (d)

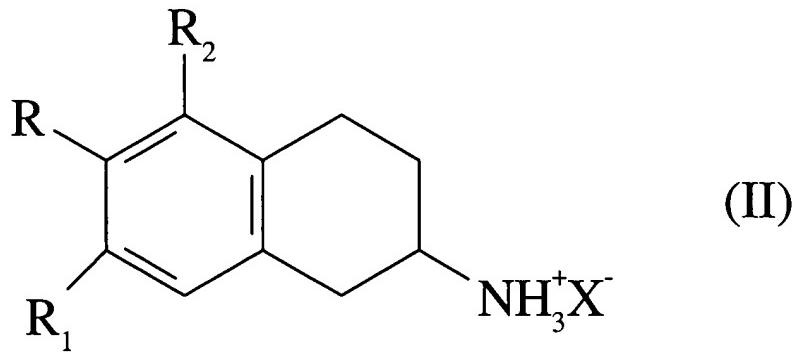
R=R₁=R₂=CH₃O,

and X⁻ is the monovalent anion of a pharmacologically acceptable acid.

9. (Previously Presented) A method of preventing or treating septic shock comprising administering to a patient in need of same an effective amount of a 2-aminotetraline of the formula (I)



or a pharmacologically acceptable salt of the formula (II)



wherein:

R and R_1 are independently halogen, hydroxy, or $\text{C}_1\text{-C}_4$ alkoxy optionally substituted in position ω with a group selected from OH , NH_2 or NR_3R_4 , wherein R_3 and

R₄ are independently H, C₁-C₄ alkyl, unsubstituted or substituted in position ω with groups OH, NH₂, C₁-C₄ alkanoyl, C₁-C₄ alkyl, carbamoyl, carbamoyloxy, amino, or amino-substituted NR₃R₄, where R₃ and R₄ have the above meanings,

R₂ is hydrogen, halogen, hydroxy or methoxy,

with the proviso that the 2-aminotetraline excludes (a) R=R₁=CH₃O or OH, R₂=H,

(b) R=F, R₁=CH₃O or OH, R₂=H, (c) R₁=-OCH₃, R=CH₃ and R₂=H, or (d)

R=R₁=R₂=CH₃O, and

X⁻ is the monovalent anion of a pharmacologically acceptable acid.

10. Canceled.

11.-12. Canceled.

13. Canceled.

14. (Previously Presented) A method of treating an inflammatory and/or autoimmune pathology induced by inflammatory cytokines, which method comprises administering to a patient in need of same an effective amount of a compound selected from the group consisting of:

S(-)-2-amino-6-fluoro-7-hydroxytetraline hydrochloride;

R(+)-2-amino-6-fluoro-7-hydroxytetraline hydrochloride;

(R,S)-2-amino-5,6-difluoro-7-methoxytetraline hydrochloride;

(R,S)-2-amino-6-fluoro-7-methyltetraline hydrochloride;

(R,S)-2-amino-7-fluoro-6-hydroxytetraline hydrochloride;

(R,S)-7-acetyl-2-amino-6-methyltetraline hydrochloride; and

(R,S)-2-amino-7-fluoro-6-methoxytetraline hydrochloride.

15. (Previously Presented) A method of preventing or treating septic shock comprising administering to a patient in need of same an effective amount of a compound selected from the group consisting of:

S(-)-2-amino-6-fluoro-7-hydroxytetraline hydrochloride;

R(+)-2-amino-6-fluoro-7-hydroxytetraline hydrochloride;

(R,S)-2-amino-5,6-difluoro-7-methoxytetraline hydrochloride;

(R,S)-2-amino-6-fluoro-7-methyltetraline hydrochloride;

(R,S)-2-amino-7-fluoro-6-hydroxytetraline hydrochloride;

(R,S)-7-acetyl-2-amino-6-methyltetraline hydrochloride; and

(R,S)-2-amino-7-fluoro-6-methoxytetraline hydrochloride.

16. Canceled.